

Produktinformation



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Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

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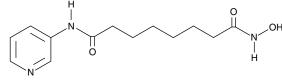
Product Information



Pyroxamide

Item No. 13870

CAS Registry No.:	382180-17-8	
Formal Name:	N ¹ -hydroxy-N ⁸ -3-pyridinyl-	H
	octanediamide	\sim $N_{\rm N}$
MF:	$C_{13}H_{19}N_{3}O_{3}$	
FW:	265.3	
Purity:	≥95%	N
Stability:	≥2 years at -20°C	
Supplied as:	A crystalline solid	
UV/Vis.:	λ _{max} : 203, 241, 279 nm	



Laboratory Procedures

For long term storage, we suggest that pyroxamide be stored as supplied at -20°C. It should be stable for at least two years.

Pyroxamide is supplied as a crystalline solid. A stock solution may be made by dissolving the pyroxamide in the solvent of choice. Pyroxamide is soluble in organic solvents such as DMSO and dimethyl formamide, which should be purged with an inert gas. The solubility of pyroxamide in these solvents is approximately 5 and 2 mg/ml, respectively.

Pyroxamide is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, pyroxamide should first be dissolved in DMSO and then diluted with the aqueous buffer of choice. Pyroxamide has a solubility of approximately 0.5 mg/ml in a 1:1 solution of DMSO:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Pyroxamide is an inhibitor of histone deacetylases (HDACs), including HDAC1 (IC₅₀ = 0.1-0.2 μ M).^{1,2} It induces growth suppression and cell death of certain types of cancer cells in culture.^{1,3}

References

- 1. Butler, L.M., Webb, Y., Agus, D.B., et al. Inhibition of transformed cell growth and induction of cellular differentiation by pyroxamide, an inhibitor of histone deacetylase. Clin. Cancer Res. 7, 962-970 (2001).
- 2. Remiszewski, S.W., Sambucetti, L.C., Atadja, P., et al. Inhibitors of human histone deacetylase: Synthesis and enzyme and cellular activity of straight chain hydroxamates. J. Med. Chem. 45(4), 753-757 (2002).
- 3. Kutko, M.C., Glick, R.D., Butler, L.M., et al. Histone deacetylase inhibitors induce growth suppression and cell death in human rhabdomyosarcoma in vitro. Clin. Cancer Res. 9, 5749-5755 (2003).

Related Products

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WARNING: This product is for laboratory research only: not for administration to humans. Not for human or veterinary DIAGNOSTIC OR THERAPEUTIC USE.

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